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U.S. PATENT DOCUMENTS

EXAMINER INITIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE (IF APPROPRIATE)

FOREIGN PATENT DOCUMENTS

DOCUMENT NO.	DATE OF PUBLICATION	COUNTRY	CLASS	SUBCLASS	TRANSLATION YES	NO

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

AA	✓	Ahmad et al., "Alopecia Universalis Associated with a Mutation in the Human <i>hairless</i> Gene," Science, 1998, Vol. 279, p. 720-724
AB	✓	Armstrong et al., "Haploinsufficiency of desmoplakin causes a striate subtype of palmoplantar keratoderma", Human Molecular Genetics, 1999, Vol. 8, No.1, p. 143-148
AC	✓	Dong et al., "Frequent Somatic Mutations in <i>Serine/Threonine Kinase 11</i> /Peutz-Jeghers Syndrome Gene in Left-sided Colon Cancer," Cancer Research, 1998, Vol. 58, p. 3787-3790
AD	✓	Frank et al., "Exposing the human <i>nude</i> phenotype," Nature, 1999, Vol. 398, p. 473-474
AE	✓	Korge et al., "A Mutational Hotspot in the 2B Domain of Human Hair Basic Keratin 6 (hHb6) in Monilethrix Patients," 1998, Vol. 111, No. 5, p 896-898
AF	✓	Marsh et al., "Germline mutations in <i>PTEN</i> are present in Bannayan-Zonana syndrome," Nature Genetics, 1997, Vol. 16, p. 333-334
AG	✓	McGrath et al., "Mutations in the plakophilin 1 gene result in ectodermal dysplasia/skin fragility syndrome," Nature Genetics, 1997, Vol. 17, p. 240-244
AH	✓	Richard et al., "Functional defects of Cx26 resulting from a heterozygous missense mutation in a family with dominant deaf-mutism and palmoplantar keratoderma," Hum Genet, 1998, Vol. 103, p. 393-399
AI	✓	Richard et al., "Mutations in the human connexin gene <i>GJB3</i> cause erythrokeratoderma variabilis," Nature Genetics, 1998, Vol. 20, p. 366-369
AJ	✓	Rickman et al., "N-terminal deletion in a desmosomal cadherin causes the autosomal dominant skin disease striate palmoplantar keratoderma," Human Molecular Genetics, 1999, Vol. 8, No. 6, p. 971-976
AK	✓	Rowan et al., "Somatic Mutations in the Peutz-Jeghers (LKB1/STKII) Gene in Sporadic Malignant Melanomas," The Journal of Investigative Dermatology, 1999, Vol. 112, No. 4, p. 509-511
AL	✓	Sakuntabhai et al., "Mutations in <i>ATP2A2</i> , encoding a Ca ²⁺ pump, cause Darier disease," Nature Genetics, 1999, Vol. 21, p. 271-277
AM	✓	Winter et al., "Mutations in the hair cortex keratin hHb6 cause the inherited hair disease monilethrix," Nature Genetics, 1997, Vol. 16, p. 372-374

EXAMINER

Joe Wontach

DATE CONSIDERED

11/28/01

EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through Citation if not in conformance and not considered. Include copy of this form with next communication to applicant.